

**SA 924.** Mr. TOOMEY submitted an amendment intended to be proposed to amendment SA 891 proposed by Mr. SCHUMER to provide for reconciliation pursuant to title II of S. Con. Res. 5; which was ordered to lie on the table; as follows:

In section 2301, add at the end the following:

“(d) **EXPERTISE REQUIREMENT.**—Any amount awarded to a State, local, Tribal, or territorial public health department pursuant to subsection (b)(2) shall be conditioned on such public health department agreeing to make such award funds available—

“(1) only to entities with which the public health department has an established relationship, and based on demonstrated expertise of such entities in vaccine distribution and administration; and

“(2) with special consideration given to such entities serving medically underserved areas.”.

**SA 925.** Mr. TOOMEY submitted an amendment intended to be proposed to amendment SA 891 proposed by Mr. SCHUMER to the bill H.R. 1319, to provide for reconciliation pursuant to title II of S. Con. Res. 5; which was ordered to lie on the table; as follows:

Strike section 9662.

**SA 926.** Mr. MARSHALL submitted an amendment intended to be proposed to amendment SA 891 proposed by Mr. SCHUMER to the bill H.R. 1319, to provide for reconciliation pursuant to title II of S. Con. Res. 5; which was ordered to lie on the table; as follows:

At the end of title II, insert the following:

**Subtitle M—Food and Drug Administration**

**SEC. 2931. USING EMERGENCY USE AUTHORIZATION DATA AND REAL WORLD EVIDENCE GATHERED DURING AN EMERGENCY TO SUPPORT DRUG, BIOLOGICAL PRODUCT, AND PRE-MARKET DEVICE APPLICATIONS.**

(a) **IN GENERAL.**—Data generated to support an authorization under section 564 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360bbb-3) with respect to a drug, biological product, or device, and real world evidence relating to such drug, biological product, or device used pursuant to such authorization, may constitute valid scientific evidence, and shall be considered for purposes of—

(1) reviewing submissions pursuant to section 505 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 355) and section 351 of the Public Health Service Act (42 U.S.C. 262);

(2) reviewing submissions pursuant to sections 510(k), 513(f), and 515 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 21 U.S.C. 360(k), 360c(f), or 360e); and

(3) otherwise meeting the requirements of such Act and such section 351 of the Public Health Service Act.

(b) **APPLICABILITY OF CERTAIN CATEGORIZATIONS FOR PREMARKET DEVICE REVIEW.**—In the case of a device receiving an authorization under section 564 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360bbb-3) for which the Secretary has determined, in accordance with subsection (m) of such section, that a laboratory examination or procedure associated with such device is deemed to be in the category of examinations and procedures described in section 353(d)(3) of the Public Health Service Act (42 U.S.C. 262), such determination shall apply with regard to a submission pursuant to section 510(k), 513(f), or 515 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 21 U.S.C. 360(k),

360c(f), or 360e) for such device, unless the Secretary (taking into account any applicable conditions specified pursuant to subsection (m)(2) of section 564 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360bbb-3)) identifies new information not included in the request for authorization that indicates that the criteria under section 353(d)(3) of the Public Health Service Act (42 U.S.C. 262) are not met.

(c) **RULE OF CONSTRUCTION.**—Nothing in this section shall be construed as altering the review standards or otherwise affecting the requirements under section 505, 510(k), 513(f), or 515 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 21 U.S.C. 355, 360(k), 360c(f), or 360e) or under section 351 of the Public Health Service Act (42 U.S.C. 262) for the clearance or approval of a device, approval of a drug, or licensure of a biological product.

**SA 927.** Mr. MARSHALL submitted an amendment intended to be proposed to amendment SA 891 proposed by Mr. SCHUMER to the bill H.R. 1319, to provide for reconciliation pursuant to title II of S. Con. Res. 5; which was ordered to lie on the table; as follows:

Strike section 9301 and insert the following:

**SEC. 9301. ADDITIONAL FUNDING FOR AGING AND DISABILITY SERVICES PROGRAMS.**

Subtitle A of title XX of the Social Security Act (42 U.S.C. 1397-1397h) is amended by adding at the end the following:

**“SEC. 2010. ADDITIONAL FUNDING FOR AGING AND DISABILITY SERVICES PROGRAMS.**

“(a) **APPROPRIATION.**—In addition to amounts otherwise available, there is appropriated for fiscal year 2021, out of any money in the Treasury not otherwise appropriated, \$276,000,000, to remain available until expended, to carry out the programs described in subtitle B.

“(b) **USE OF FUNDS.**—Subject to subsection (c), of the amounts made available by subsection (a)—

“(1) \$88,000,000 shall be made available to carry out the programs described in subtitle B in fiscal year 2021, of which not less than an amount equal to \$100,000,000 minus the amount previously provided in fiscal year 2021 to carry out section 2042(b) shall be made available to carry out such section; and

“(2) \$188,000,000 shall be made available to carry out the programs described in subtitle B in fiscal year 2022, of which not less than \$100,000,000 shall be for activities described in section 2042(b).

“(c) **LIMITATION ON USE OF FUNDS.**—None of the amounts made available by subsection (a) may be paid, obligated, or otherwise expended to carry out the programs described in subtitle B in a State that does not have COVID-19 medical liability protections for any health care provider who works in a long term care facility or nursing facility (as such terms are defined in section 2011).”.

**SA 928.** Mr. MARSHALL submitted an amendment intended to be proposed to amendment SA 891 proposed by Mr. SCHUMER to the bill H.R. 1319, to provide for reconciliation pursuant to title II of S. Con. Res. 5; which was ordered to lie on the table; as follows:

On page 278, line 22, strike the period and insert “; and \$100,000,000 shall be available to purchase medical supplies that are made in the United States.”.

**SA 929.** Mr. MARSHALL submitted an amendment intended to be proposed

to amendment SA 891 proposed by Mr. SCHUMER to the bill H.R. 1319, to provide for reconciliation pursuant to title II of S. Con. Res. 5; which was ordered to lie on the table; as follows:

At the end of title V, add the following:

**SEC. 5007. PROHIBITION ON PPP LOANS FOR ABORTION PROVIDERS.**

(a) **IN GENERAL.**—Section 7(a)(36) of the Small Business Act (15 U.S.C. 636(a)(36)) is amended by adding at the end the following:

“(T) **PROHIBITION ON COVERED LOANS FOR ABORTION PROVIDERS.**—

“(i) **IN GENERAL.**—Except as provided in clause (ii), no individual or entity that provides abortions shall be eligible to receive a covered loan.

“(ii) **EXCEPTIONS.**—Clause (i) shall not apply to—

“(I) a hospital, as defined in section 1861(e) of the Social Security Act (42 U.S.C. 1395x(e)); or

“(II) an entity that exclusively provides abortions described in section 507(a) of the Further Consolidated Appropriations Act, 2020 (Public Law 116-94).”.

(b) **EFFECTIVE DATE.**—The amendment made by this section shall be effective as if included in the enactment of the CARES Act (Public Law 116-136).

(c) **INSPECTOR GENERAL REPORT.**—Not later than 6 months after the date of enactment of this Act, the Inspector General of the Small Business Administration shall conduct an investigation and submit to Congress a report on the number of loans made to the Planned Parenthood Federation of America pursuant to section 7(a)(36) of the Small Business Act (15 U.S.C. 636(a)(36)) and to other individuals or entities that provide abortions.

**SA 930.** Mr. MARSHALL (for himself, Mr. KENNEDY, Mrs. BLACKBURN, Mr. BRAUN, Mr. ROMNEY, Mr. YOUNG, Mr. GRASSLEY, and Mr. COTTON) submitted an amendment intended to be proposed by him to the bill H.R. 1319, to provide for reconciliation pursuant to title II of S. Con. Res. 5; which was ordered to lie on the table; as follows:

In section 2003, add at the end the following:

(8) an institution shall not be eligible to receive an allocation under this section unless, not later than 60 days after the date of enactment of this Act, that institution certifies to the Secretary of Education that the institution does not have a partnership in effect with a cultural institute directly or indirectly funded by the Government of the People's Republic of China (referred to as a “Confucius Institute”).

**SA 931.** Mr. MARSHALL submitted an amendment intended to be proposed to amendment SA 891 proposed by Mr. SCHUMER to the bill H.R. 1319, to provide for reconciliation pursuant to title II of S. Con. Res. 5; which was ordered to lie on the table; as follows:

At the end of title VII, add the following:

**Subtitle G—Limitation on Use of Funds**

**SEC. 7701. RELIEF FUND FOR CERTAIN PIPELINE WORKERS.**

None of the funds provided by this title may be expended until a relief fund is established to compensate individuals who have lost employment due to the cancellation of the Keystone XL Pipeline pursuant to section 6 of Executive Order 13990 (86 Fed. Reg. 7041 (January 25, 2021)), which revoked the Presidential Permit of March 29, 2019 (84 Fed. Reg. 13101 (April 3, 2019)) authorizing Trans-Canada Keystone Pipeline, L.P., to construct, connect, operate, and maintain pipeline facilities at the international border of the United States and Canada.